Identifiers
1. Enter Refid
2. First author last name
3. Year of
4. Reviewer initials
5. Study Question
☐ KQ1: Preoperative Evaluation
KQ2: Prehabilitation
KQ3: Regional versus General
KQ4: Intravenous versus Inhaled
KQ5: Potentially Inappropriate Medications
KQ6: Pharmacologic Delirium Prophylaxis
KQ7: Postoperative Regional Anesthesia
KQ8: PACU Delirium Screening
6. Start date (MM/YY)
7. End date (MM/YY)
Q. Desistand
8. Registered ——————————————————————————————————
○ No
○ Uncertain
Clear Response ()
9. NCT or other registration number
10. Country
Select an Answer V
11. Multinational (list)
12. Non-very high HDI country < 0.8 (see WHO report (http://hdr.undp.org/en/content/latest-human-development-index-ranking))
Yes
Clear Response ()
Funding
13. Sponsor
O Public or nonprofit
○ Industry
O Public, nonprofit, and industry
○ None
○ Not reported
Clear Response ()
14. Sponsor description

15. Author COI
○ No COI
○ Report COI
○ Not reported
Clear Response ()
Characteristics
17. Study design
Randomized Controlled Trial (parallel)
○ Cluster Randomized
○ Crossover Trial
○ Non-randomized clinical trial
○ Before-after/Interrupted time series
Prospective Cohort (observational)
Retrospective Cohort (observational)
○ Cross-sectional
○ Case-control
○ Fully paired (diagnostic)
○ Case series
Other
19. Research letter
Yes
Clear Response ()
20. Pilot study
Yes
Clear Response ()
21. Trial Phase
Phase I
O Phase II
O Phase III
○ Phase IV
Clear Response ()
22. Setting description
23. Setting Ambulatory
Hospital
Other
Not stated

25. Inclusion criteria (short description)
26. Exclusion criteria (short description)
20. Exclusion criteria (short description)
27. Number of centers
28. Subgroups Obesity Obiabetes Age Osex Clear Response ()
Outcomes (see registration if applicable: NCT (https://clinicaltrials.gov/), ICTRP (https://trialsearch.who.int/Default.aspx), UMIN Japan (https://center6.umin.ac.jp/cgi-open-bin/ctr_e/index.cgi? function=02))
29. Primary outcome(s)
Postoperative delirium Delayed neurocognitive recovery Stroke Intraoperative awareness Quality of recovery Depression Patient/caregiver satisfaction
Valued life activities Functional status (ADL, IADL, mobility) HRQoL Pain Opioid use Complications Length of stay Discharge location Readmission
Mortality (30-, 180-, 365-day) Other
31. Secondary outcomes
Postoperative delirium Delayed neurocognitive recovery Stroke Intraoperative awareness Quality of recovery Depression Patient/caregiver satisfaction Valued life activities Functional status (ADL, IADL, mobility) HRQoL Pain Opioid use Complications Length of stay Discharge location Readmission
Mortality (30-, 180-, 365-day) Other
33. Other outcomes not specified as primary or secondary (eg, as recorded in NCT registry)
Postoperative delirium Delayed neurocognitive recovery Stroke Intraoperative awareness Quality of recovery Depression Patient/caregiver satisfaction
Valued life activities Functional status (ADL, IADL, mobility) HRQoL Pain Opioid use Complications Length of stay Discharge location Readmission
Mortality (30-, 180-, 365-day) Other
35. Source for outcomes
Publication
Registry
Protocol
36. Arms
37. Anesthetic
General
Regional
Sedation
□ Not reported
38. Type of surgery (if none state "None")
Various (4 or more)
Abdominal
☐ Cardiac
☐ Colorectal
Gynecological
Gastrointestinal
General

Head and neck		
Hepatic		
Neurosurgical		
Opthalmic		
Oral/Maxillofacial		
Orthopedic, total knee arthroplasty		
Orthopedic, total hip arthroplasty		
Orthopedic, hip fracture		
Orthopedic, other or unspecified		
Otolaryngological (ENT)		
Plastic/Cosmetic		
Spine		
Thoracic		
Urologic		
Vascular		
Other		
39. List of specific surgical procedures so	separated by semicolons	
43. Number randomized		
44. Number analyzed		
45. Notes		
46. Main study results		
47. Comment		
Tr. Comment		
47. Comment		
7. Comment		
7. Comment		
7). Comment		